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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/649,193	08/26/2003	Marilyn H. Perrin	SALK1740-10 (088802-3218)	5260	
30542 FOLEY & LA	7590 05/16/2007 RDNER LLP	INER			
P.O. BOX 80278			BORGEEST, CHRISTINA M		
SAN DIEGO,	CA 92138-0278		ART UNIT PAPER NUMBER		
			1649		
			MAIL DATE	DELIVERY MODE	
			05/16/2007	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)		
	10/649,193	PERRIN ET AL.		
Office Action Summary	Examiner	Art Unit		
	Christina Borgeest	1649		
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	orrespondence address		
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period way reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tin vill apply and will expire SIX (6) MONTHS from 1, cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).		
Status		•		
1) Responsive to communication(s) filed on 28 Fe	ebruary 2007.			
,	This action is FINAL . 2b)⊠ This action is non-final.			
3) Since this application is in condition for allowar				
closed in accordance with the practice under E	Ex parte Quayle, 1935 C.D. 11, 4	53 O.G. 213.		
Disposition of Claims				
4) ⊠ Claim(s) 1,2,5-11,13,14 and 19 is/are pending 4a) Of the above claim(s) is/are withdray 5) □ Claim(s) is/are allowed. 6) ⊠ Claim(s) 1,2,5,6,8-11,13,14 and 19 is/are reject 7) ⊠ Claim(s) 7 is/are objected to. 8) □ Claim(s) are subject to restriction and/or	wn from consideration.			
Application Papers				
9) The specification is objected to by the Examine		.		
10) The drawing(s) filed on is/are: a) acc				
Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct				
11) The oath or declaration is objected to by the Ex				
Priority under 35 U.S.C. § 119				
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority document application from the International Bureau * See the attached detailed Office action for a list	s have been received. s have been received in Applicat rity documents have been receiv u (PCT Rule 17.2(a)).	ion No ed in this National Stage		
, , , , , , , , , , , , , , , , , , ,				
Attachment(s) 1) Notice of References Cited (PTO-892)	4) Interview Summary	y (PTO-413)		
 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date 	Paper No(s)/Mail D 5) Notice of Informal 6) Other:			

DETAILED ACTION

Formal Matters

The response filed 28 February 2007 is acknowledged. Claims 1 and 11 are amended. Claims 3-4, 12, 15-18 and 20 are cancelled. Claims 1-2, 5-11, 13-14 and 19 are under consideration.

The text of those sections of 35 U.S.C. not included in this action can be found in a prior office action mailed 1 May 2006.

Objections/Rejections Withdrawn

Priority

Applicant's claim for receiving the benefit of an earlier filing date under 35 U.S.C. [120] of 23 August 1993 was acknowledged but not granted by the Examiner, and the effective filing date determined by the Examiner was 12 November 1998, as set forth at p. 3 of the previous Office actions (mailed 1 May 2006 and 29 November 2006). Applicants have properly showed that they have support for the claimed subject matter in each of the parent applications dating back to 23 August 1993 at p. 8 of their arguments (see particularly the Table at p. 8 of their Arguments). Applicants have shown continuity via copendency in the chain of intervening applications and continuity of subject matter in the chain of intervening applications (MPEP 201.11), thus the priority date of 23 August 1993 is *granted*.

Claim Rejections - 35 USC § 112, second paragraph

The rejection of claim 1, and claims 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, which depend from 1, under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention is withdrawn in response to Applicants' amendment of claims 1 and 11, cancellation claims 3-4 and upon reconsideration because previously granted U.S. patents 5,728,545 and 6,638,905 contain comparable language to the claims as currently recited.

Claim Rejections - 35 USC § 112, first paragraph

The rejection of claims 1-6, 8-11, 14-20 under 35 U.S.C. 112, first paragraph, for scope of enablement is withdrawn in response to Applicants amendment of claims 1 and 11, cancellation of claims 3-4, 12, 15-18 and 20 and upon reconsideration because previously granted U.S. patents 5,728,545 and 6,638,905 contain comparable language to the claims as currently recited.

Claim Rejections - 35 USC § 102

The rejection of claims 1, 2, 3, 4, 5, 6, 7, 9, 10, 11, 12, 13, 15, 16, 17, 18, 19 and 20 under 35 U.S.C. 102(b) as being anticipated by Laurent et al. (FEBS, 1993; 335: 1-5) is withdrawn because of the cancellation of claims 3-4, 12, 15-18 and 20 and because a priority date of 23 August 1993 was granted and the publication by Laurent et al. no longer qualifies as prior art.

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The rejection of claims 1, 8 13 and 14 under 35 U.S.C. 102(b) as being anticipated by Chen et al. (Proc Natl Acad Sci. 1993; 90: 8967-8971—on IDS filed 12 November 2003) is withdrawn because a priority date of 23 August 1993 was granted and the publication by Chen et al. no longer qualifies as prior art.

Double Patenting

The rejection of claims 1-11 and 13-20 on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1-8 of U.S. Patent No. 5,728,545 (the '545 patent) is withdrawn in response to Applicants cancellation of claims 3-4, 12, 15-18 and 20 and because a restriction requirement between the DNA and protein was made, thus the rejection over the '545 patent is not proper.

Rejection Maintained/New Rejection/Objection Claim Objections

Claim 7 is objected to as being dependent upon a rejected base claim, but would be allowable if rewritten in independent form including all of the limitations of the base claim and any intervening claims.

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Claim Rejections - 35 USC § 112, first paragraph

The rejection of claim 13 for scope of enablement as set forth in the previous Office actions (mailed 1 May 2006 and 29 November 2006) is maintained for reasons of record and the following. Claim 13 recites "[a] pure polypeptide comprising at least 15 contiguous amino acids of the amino acid sequence set forth in SEQ ID NO: 15, wherein said polypeptide is about 70% pure (by weight of total proteins). The claim requires only 15 contiguous amino acids without any functional requirement, thus it would require undue experimentation on the part of the person of ordinary skill in the art to use the polypeptides encompassed by the claim. Rijkers et al. (ChemBioChem 2004; 5: 340-348) teach a number of inactive fragments of astressin that do not have biological activity (see Table I, p. 342-343 and p. 343, left column, 1st paragraph). Although not the same protein as recited in claim 13; astressin is a CRF receptor antagonist, and the principle is the same; namely that certain residues must be preserved in order to preserve structure-activity relationships. The problem of predicting protein structure from sequence data and in turn utilizing predicted structural determinations to ascertain functional aspects of the protein is extremely complex. Certain positions in the sequence are critical to the protein's structure/function relationship, e.g. such as various sites or regions directly involved in binding, activity and in providing the correct three-dimensional spatial orientation of binding and active sites. These or other regions may also be critical determinants of antigenicity. These regions can tolerate only relatively conservative substitutions or no substitutions (see Wells, 1990, Biochemistry 29:8509-8517; Ngo et al., 1994, The Protein Folding Problem

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and Tertiary Structure Prediction, pp. 492-495). However, Applicant has provided little or no guidance beyond the mere presentation of sequence data to enable one of ordinary skill in the art to determine, without undue experimentation, which fragments of SEQ ID NO: 15 have functional activity. Although the specification outlines artrecognized procedures for producing and screening for active fragments, this is not adequate guidance, but is merely an invitation to the artisan to use the current invention as a starting point for further experimentation. Even if an active or binding site were identified in the specification, they may not be sufficient, as the ordinary artisan would immediately recognize that an active or binding site must assume the proper threedimensional configuration to be active, which conformation is dependent upon surrounding residues; therefore substitution of non-essential residues can often destroy activity. The art recognizes that function cannot be predicted from structure alone (Bork, 2000, Genome Research 10:398-400; Skolnick et al., 2000, Trends in Biotech. 18(1):34-39, especially p. 36 at Box 2; Doerks et al., 1998, Trends in Genetics 14:248-250; Smith et al., 1997, Nature Biotechnology 15:1222-1223; Brenner, 1999, Trends in Genetics 15:132-133; Bork et al., 1996, Trends in Genetics 12:425-427).

Due to the large quantity of experimentation necessary to generate the large number of fragments recited in the claims and screen same for activity, the lack of direction/guidance presented in the specification regarding which structural features are required in order to provide activity, the absence of working examples directed to same, the complex nature of the invention, the state of the prior art which establishes the unpredictability of the effects of mutation on protein structure and function, and the

breadth of the claims which fail to recite any structural or functional limitations, undue experimentation would be required of the skilled artisan to make and/or use the claimed invention in its full scope.

Double Patenting

The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. A nonstatutory obviousness-type double patenting rejection is appropriate where the conflicting claims are not identical, but at least one examined application claim is not patentably distinct from the reference claim(s) because the examined application claim is either anticipated by, or would have been obvious over, the reference claim(s). See, e.g., *In re Berg*, 140 F.3d 1428, 46 USPQ2d 1226 (Fed. Cir. 1998); *In re Goodman*, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); *In re Van Ornum*, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); *In re Vogel*, 422 F.2d 438, 164 USPQ 619 (CCPA 1970); and *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) or 1.321(d) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent either is shown to be commonly owned with this application, or claims an invention made as a result of activities undertaken within the scope of a joint research agreement.

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

Claims 1-2, 5-6, 8-11, 13-14 and 19 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1-23 of U.S. Patent No. 6,638,905 (the '905 patent). Although the conflicting claims are not identical, they are not patentably distinct from each other because the instant claims as broadly recited, encompass the G protein-coupled corticotropin-releasing factor (CRF) receptor protein of the '905 patent. For example, instant claims 1, 11 (and their dependent

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claims) recite the CRF receptor protein in terms of a protein that is encoded by DNA that hybridizes to the complement of the polynucleotide sequence set forth in SEQ ID NO: 14...so as to allow identification of sequences having at least 70% nucleic acid identity with respect to SEQ ID NO: 14", and SEQ ID NO: 1 of the '905 patent has greater than 70% identity with SEQ ID NO: 14 of the instant application. For evidence of this see SCORE results reproduced below:

```
RESULT 8
AR412100
                                                        PAT 18-DEC-2003
                                1495 bp
                                         DNA
                                                 linear
LOCUS
          AR412100
          Sequence 1 from patent US 6638905.
DEFINITION
          AR412100
ACCESSION
          AR412100.1 GI:40164659
VERSION
KEYWORDS
          Unknown.
SOURCE
 ORGANISM Unknown.
          Unclassified.
             (bases 1 to 1495)
REFERENCE
          Perrin, M.H., Chen, R., Lewis, K.A., Vale, W.W. Jr., Donaldson, C.J. and
 AUTHORS
          Sawchenko, P.
          Cloning and recombinant production of CFR receptor(s)
 TITLE
          Patent: US 6638905-A 1 28-OCT-2003;
 JOURNAL
           The Salk Institute for Biological Studies; La Jolla, CA
                  Location/Qualifiers
FEATURES
                  1. .1495
    source
                   /organism="unknown"
                   /mol type="genomic DNA"
ORIGIN
                              Score 1398; DB 6;
                                                Length 1495;
  Query Match
                       88.4%;
                              Pred. No. 1e-252;
  Best Local Similarity
                       94.5%;
                                Mismatches
                                                 Indels
                                                         87;
                                                                    1;
 Matches 1495; Conservative
                             0;
                                             0;
           1 CGAGCCCGCAGCCGCCGCCGGTTCCTCTGGGATGTCCGTAGGACCCGGGCATTCAGGAC 60
Qу
             1 CGAGCCCGCAGCCGCCGGCTTCCTCTGGGATGTCCGTAGGACCCGGGCATTCAGGAC 60
Db
          61 GGTAGCCGAGCGAGCCCGAGGATGGGAGGGCACCCGCAGCTCCGTCTCAAGGCCCTT 120
Qу
             61 GGTAGCCGAGCGCGAGGATGGGAGGCACCCGCAGCTCCGTCTCGTCAAGGCCCTT 120
Db
         121 CTCCTTCTGGGGCTGAACCCCGTCTCTGCCTCCAGGACCAGCACTGCGAGAGCCTG 180
Qу
             121 CTCCTTCTGGGGCTGAACCCCGTCTCTGCCTCCCTCCAGGACCAGCACTGCGAGAGCCTG 180
Db
         181 TCCCTGGCCAGCAACATCTCAGGACTGCAGTGCAACGCATCCGTGGACCTCATTGGCACC 240
Qу
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Db	181		240
Qy	241	TGCTGGCCCGCAGCCCTGCGGGGCAGCTAGTGGTTCGGCCCTGCCCTTTTTCTAT	300
Db	241	TGCTGGCCCGCAGCCCTGCGGGGCAGCTAGTGGTTCGGCCCTGCCTTTTTCTAT	300
Qу	301	GGTGTCCGCTACAATACCACAAACAATGGCTACCGGGAGTGCCTGGCCAATGGCAGCTGG	360
Db	301	GGTGTCCGCTACAATACCACAAACAATGGCTACCGGGAGTGCCTGGCCAATGGCAGCTGG	360
Qy	361	GCCGCCCGCGTGAATTACTCCGAGTGCCAGGAGATCCTCAATGAGGAGAAAAAAAGCAAG	420
Db	361	GCCGCCGCGTGAATTACTCCGAGTGCCAGGAGATCCTCAATGAGGAGAAAAAAAGCAAG	420
Qу		GTGCACTACCATGTCGCAGTCATCATCAACTACCTGGGCCACTGTATCTCCCTGGTGGCC	480
Db	421		480
Qy		CTCCTGGTGGCCTTTGTCCTCTTTCTGCGGCTCAGGCCAGGCTGCACCCATTGGGGTGAC	
Db		CTCCTGGTGGCCTTTGTCCTCTTTCTGCGGCTC	
Qy		CAGGCAGATGGAGCCCTGGAGGTGGGGGCTCCATGGAGTGGTGCCCCATTTCAGGTTCGA	600
Db			513
Qу		AGGAGCATCCGGTGCCTGCGAAACATCATCCACTGGAACCTCATCTCCGCCTTCATCCTG	
Db		AGGAGCATCCGGTGCCTGCGAAACATCATCCACTGGAACCTCATCTCCGCCTTCATCCTG	
Qу		CGCAACGCCACCTGGTTCGTGGTCCAGCTAACCATGAGCCCCGAGGTCCACCAGAGCAAC	
Db		CGCAACGCCACCTGGTTCGTGGTCCAGCTAACCATGAGCCCCGAGGTCCACCAGAGCAAC	
Qу			780
Db		GTGGGCTGGTGCAGGTTGGTGACAGCCGCCTACAACTACTTCCATGTGACCAACTTCTTC	
Qy		TGGATGTTCGGCGAGGGCTGCTACCTGCACACAGCCATCGTGCTCACCTACTCCACTGAC	
Db		TGGATGTTCGGCGAGGGCTGCTACCTGCACACAGCCATCGTGCTCACCTACTCCACTGAC	
Qy		CGGCTGCGCAAATGGATGTTCATCTGCATTGGCTGGGGTGTGCCCTTCCCCATCATTGTG	
Db		CGGCTGCGCAAATGGATGTTCATCTGCATTGGCTGGGGTGTGCCCTTCCCCATCATTGTG GCCTGGGCCATTGGGAAGCTGTACTACGACAATGAGAAGTGCTGGTTTGGCAAAAGGCCT	
Qy Db		GCCTGGGCCATTGGGAAGCTGTACTACGACAATGAGAAGTGCTGGTTTGGCAAAAGGCCT	
Db		GGGGTGTACACCGACTACATCTACCAGGGCCCCATGATCCTGGTCCTGATCAATTTC	
Qy Db		GGGGTGTACACCGACTACATCTACCAGGGCCCCATGATCCTGGTCCTGATCATTTC	
D.1.	0 / 4		

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```
1021 ATCTTCCTTTTCAACATCGTCCGCATCCTCATGACCAAGCTCCGGGCATCCACCACGTCT 1080
Qу
         934 ATCTTCCTTTTCAACATCGTCCGCATCCTCATGACCAAGCTCCGGGCATCCACCACGTCT 993
Db
     1081 GAGACCATTCAGTACAGGAAGGCTGTGAAAGCCACTCTGGTGCTGCTGCCCCTCCTGGGC 1140
Qу
         994 GAGACCATTCAGTACAGGAAGGCTGTGAAAGCCACTCTGGTGCTGCTGCCCCTCCTGGGC 1053
Db
     1141 ATCACCTACATGCTGTTCTTCGTCAATCCCGGGGAGGATGAGGTCTCCCGGGTCGTCTTC 1200.
Qу
         1054 ATCACCTACATGCTGTTCTTCGTCAATCCCGGGGAGGATGAGGTCTCCCGGGTCGTCTTC 1113
Db
     1201 ATCTACTTCAACTCCTTGCTGGAATCCTTCCAGGGCTTCTTTGTGTCTGTGTTCTACTGT 1260
Qу
         1114 ATCTACTTCAACTCCTTGGAATCCTTCCAGGGCTTCTTTGTGTCTGTGTTCTACTGT 1173
Db
     1261 TTCCTCAATAGTGAGGTCCGTTCTGCCATCCGGAAGAGGTGGCACCGGTGGCAGGACAAG 1320
Qу
         1174 TTCCTCAATAGTGAGGTCCGTTCTGCCATCCGGAAGAGGTGGCACCGGTGGCAGGACAAG 1233
Db
     1321 CACTCGATCCGTGCCCGAGTGGCCCGTGCCATCCCCACCCTCCCCAACCCGTGTC 1380
Qу
         1234 CACTCGATCCGTGCCCGAGTGGCCCGTGCCATGTCCATCCCCACCTCCCCAACCCGTGTC 1293
Db
     1381 AGCTTTCACAGCATCAAGCAGTCCACAGCAGTCTGAGCTGGCAGGTCATGGAGCAGCCCC 1440
Qу
         1294 AGCTTTCACAGCATCAAGCAGTCCACAGCAGTCTGAGCTGGCAGGTCATGGAGCAGCCCC 1353
Db
     Qу
         Db
     1501 GACCTGTTAGGTCTCATGCCCACTCCCCCAGGAGCAGCTGGCACTGACAGCCTGGGGGGG 1560
Qу
         1414 GACCTGTTAGGTCTCATGCCCACTCCCCCAGGAGCAGCTGGCACTGACAGCCTGGGGGGG 1473
Db
     1561 CCGCTCTCCCCCTGCAGCCGTG 1582
Qу
         1474 CCGCTCTCCCCCTGCAGCCGTG 1495
Db
```

In addition, the CRF receptor protein recited in the claims of the '905 patent has greater than 70% identity with SEQ ID NO: 15. For evidence of this see SCORE results reproduced below

RESULT 8

US-09-191-724-1

[;] Sequence 1, Application US/09191724

[;] Patent No. 6638905

```
; GENERAL INFORMATION:
 APPLICANT: Perrin, Marilyn H.
 APPLICANT: Chen, Ruoping
 APPLICANT: Lewis, Kathy A.
 APPLICANT: Vale Jr., Wylie W.
 APPLICANT: Donaldson, Cynthia J.
 APPLICANT: Sawchenko, Paul
 TITLE OF INVENTION: Cloning and Recombinant Production of
 TITLE OF INVENTION: CRF Receptor(s)
 FILE REFERENCE: Salk1748
 CURRENT APPLICATION NUMBER: US/09/191,724
 CURRENT FILING DATE: 1998-11-12
 EARLIER APPLICATION NUMBER: US 08/374,009
 EARLIER FILING DATE: 1995-01-17
 EARLIER APPLICATION NUMBER: US 08/353,537
  EARLIER FILING DATE: 1994-12-09
  EARLIER APPLICATION NUMBER: PCT/US94/05908
 EARLIER FILING DATE: 1993-05-25
; EARLIER APPLICATION NUMBER: US 08/110,286
; EARLIER FILING DATE: 1993-08-23
 EARLIER APPLICATION NUMBER: US 08/079,320
  EARLIER FILING DATE: 1993-06-18
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 1
  LENGTH: 1495
   TYPE: DNA
   ORGANISM: Homo sapiens
  FEATURE:
  NAME/KEY: CDS
  LOCATION: (82)...(1326)
   OTHER INFORMATION: /product = "Human pituitary CRF-receptor"
   OTHER INFORMATION: /note= "This sequence is encoded by clone
   OTHER INFORMATION: "CRF-R1"."
US-09-191-724-1
                       88.4%; Score 1398; DB 3; Length 1495;
 Query Match
 Best Local Similarity 94.5%; Pred. No. 0;
 Matches 1495; Conservative 0; Mismatches
                                             0; Indels 87; Gaps
1;
          1 CGAGCCCGCAGCCGCCGCCGGTTCCTCTGGGATGTCCGTAGGACCCGGGCATTCAGGAC 60
            1 CGAGCCCGCAGCCGCCCGCTTCCTCTGGGATGTCCGTAGGACCCGGGCATTCAGGAC 60
          61 GGTAGCCGAGCGCGAGGATGGGAGGGCACCCGCAGCTCCGTCTCGTCAAGGCCCTT 120
Qу
            61 GGTAGCCGAGCGCGAGGATGGGAGGGCACCCGCAGCTCCGTCTCGTCAAGGCCCTT 120
Db
         121 CTCCTTCTGGGGCTGAACCCCGTCTCTGCCTCCCTCCAGGACCAGCACTGCGAGAGCCTG 180
Qу
            121 CTCCTTCTGGGGCTGAACCCCGTCTCTGCCTCCCAGGACCAGCACTGCGAGAGCCTG 180
Db
         181 TCCCTGGCCAGCAACATCTCAGGACTGCAGTGCAACGCATCCGTGGACCTCATTGGCACC 240
Qу
```

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Db	181	TCCCTGGCCAGCAACATCTCAGGACTGCAGTGCAACGCATCCGTGGACCTCATTGGCACC	240
Qy	241	TGCTGGCCCGCAGCCCTGCGGGGCAGCTAGTGGTTCGGCCCTGCCCTTTTTCTAT	300
Db	241	TGCTGGCCCCGCAGCCCTGCGGGGCAGCTAGTGGTTCGGCCCTGCCCTTTTTCTAT	300
Qy	301	GGTGTCCGCTACAATACCACAAACAATGGCTACCGGGAGTGCCTGGCCAATGGCAGCTGG	360
Db	301	GGTGTCCGCTACAATACCACAAACAATGGCTACCGGGAGTGCCTGGCCAATGGCAGCTGG	360
Qy	361	GCCGCCCGCGTGAATTACTCCGAGTGCCAGGAGATCCTCAATGAGGAGAAAAAAAGCAAG	.420
Db	361	GCCGCCGCGTGAATTACTCCGAGTGCCAGGAGATCCTCAATGAGGAGAAAAAAAA	420
Qy	421	GTGCACTACCATGTCGCAGTCATCATCAACTACCTGGGCCACTGTATCTCCCTGGTGGCC	48Ö
Db	421		480
Qy	481	CTCCTGGTGGCCTTTGTCCTCTTTCTGCGGCTCAGGCCAGGCTGCACCCATTGGGGTGAC	540
Db	481		513
Qy	541	CAGGCAGATGGAGCCCTGGAGGTGGGGGCTCCATGGAGTGGTGCCCCATTTCAGGTTCGA	600
Db	514		513
Qy	601	AGGAGCATCCGGTGCCTGCGAAACATCATCCACTGGAACCTCATCTCCGCCTTCATCCTG	660
Db	514		573
Qy	661	CGCAACGCCACCTGGTTCGTGGTCCAGCTAACCATGAGCCCCGAGGTCCACCAGAGCAAC	720
Db	574		633
Qy	721	GTGGGCTGGTGCAGGTTGGTGACAGCCGCCTACAACTACTTCCATGTGACCAACTTCTTC	780
Db	634		693
Qy	781	TGGATGTTCGGCGAGGGCTGCTACCTGCACACAGCCATCGTGCTCACCTACTCCACTGAC	840
Db	694		753
Qу	841	CGGCTGCGCAAATGGATGTTCATCTGCATTGGCTGGGGTGTGCCCTTCCCCATCATTGTG	900
Db	754		813
Qу	901	GCCTGGGCCATTGGGAAGCTGTACTACGACAATGAGAAGTGCTGGTTTGGCAAAAGGCCT	960
Db	814		873
Qy	961	GGGGTGTACACCGÁCTACATCTACCAGGGCCCCATGATCCTGGTCCTGATCAATTTC	102
Db	874		933

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Qy ·	1021	ATCTTCCTTTTCAACATCGTCCGCATCCTCATGACCAAGCTCCGGGCATCCACCACGTCT	1080
Db	934	ATCTTCCTTTTCAACATCGTCCGCATCCTCATGACCAAGCTCCGGGCATCCACCACGTCT	993
Qy	1081	GAGACCATTCAGTACAGGAAGGCTGTGAAAGCCACTCTGGTGCTGCCCCTCCTGGGC	1140
Db	994	GAGACCATTCAGTACAGGAAGGCTGTGAAAGCCACTCTGGTGCTGCTGCCCCTCCTGGGC	1053
Qy	1141	ATCACCTACATGCTGTTCTTCGTCAATCCCGGGGAGGATGAGGTCTCCCGGGTCGTCTTC	1200
Db	1054	ATCACCTACATGCTGTTCTTCGTCAATCCCGGGGAGGATGAGGTCTCCCGGGTCGTCTTC	1113
Qy	1201	ATCTACTTCAACTCCTTCCTGGAATCCTTCCAGGGCTTCTTTGTGTCTGTGTTCTACTGT	1260
Db	1114	ATCTACTTCAACTCCTTGCAGAATCCTTCCAGGGCTTCTTTGTGTCTGTGTTCTACTGT	1173
Qy	1261	TTCCTCAATAGTGAGGTCCGTTCTGCCATCCGGAAGAGGTGGCACCGGTGGCAGGACAAG	1320
Db	1174	TTCCTCAATAGTGAGGTCCGTTCTGCCATCCGGAAGAGGTGGCACCGGTGGCAGACAAG	1233
Qy	1321	CACTCGATCCGTGCCCGAGTGGCCCGTGCCATCTCCCCACCTCCCCAACCCGTGTC	1380
Db	1234	CACTCGATCCGTGCCCGAGTGGCCCGTGCCATGTCCATCCCCAACCCGTGTC	1293
Qy	1381	AGCTTTCACAGCATCAAGCAGTCCACAGCAGTCTGAGCTGGCAGGTCATGGAGCAGCCCC	1440
Db	1294	AGCTTTCACAGCATCAAGCAGTCCACAGCAGTCTGAGCTGGCAGGTCATGGAGCCCC	1353
Qу	1441	CAAAGAGCTGTGGCTGGGGGGATGACGGCCAGGCTCCCTGACCACCCTGCCTG	1500
Db	1354		1413
Qy	1501	GACCTGTTAGGTCTCATGCCCACTCCCCCAGGAGCAGCTGGCACTGACAGCCTGGGGGGG	1560
Db	1414	GACCTGTTAGGTCTCATGCCCACTCCCCCAGGAGCAGCTGGCACTGACAGCCTGGGGGGG	1473
Qу	1561	CCGCTCTCCCCTGCAGCCGTG 1582	
Db	1474		

In addition, claim 5 (80% nucleic acid identity with respect to SEQ ID NO: 14) and claim 6 (90% nucleic acid identity with respect to SEQ ID NO: 14), are encompassed by the claims of the '905 patent since SEQ ID NO: 1 (recited in the patented claims) has over 94% similarity with SEQ ID NO: 14 (recited in the instant claims). The instant claims are very broad, and recite the claimed CRF receptor protein

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in terms of hybridization to the complement of the DNA encoding the protein, thus encompass the claims of the '905 patent.

Conclusion

Claims 1-2, 5-6, 8-11, 13-14 and 19 are rejected. Claim 7 contains allowable subject matter.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Christina Borgeest whose telephone number is 571-272-4482. The examiner can normally be reached on 8:00-4:30.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Janet Andres, Ph.D. can be reached on 571-272-0867. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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